

Supporting Information

Turer et al. 10.1073/pnas.1621400114

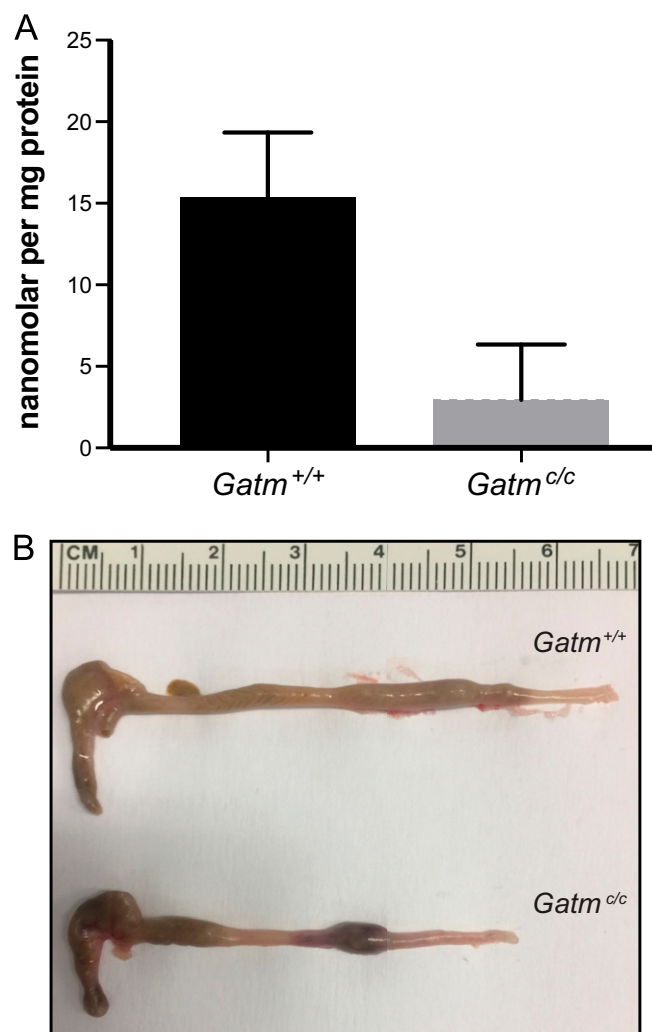


Fig. S1. Creatine content and colitis induced shortening of *Gatm*^{cl^c} colons. (A) Creatine concentration in *Gatm*^{cl^c} ($n = 4$) and littermate control (*Gatm*^{+/+}, $n = 4$) colonic epithelial cells. $P < 0.005$. (B) Colons of *Gatm*^{cl^c} mice exposed to 1.4% DSS for 8 d exhibited shortening. Representative colons are shown for each genotype.

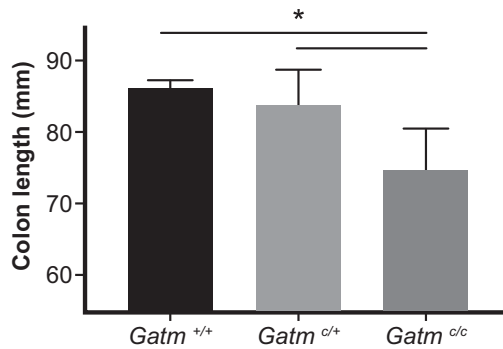


Fig. S2. Colons of trinitrobenzenesulfonic acid (TNBS)-treated *Gatm*^{dc} mice. Colons of *Gatm*^{dc} displayed shortening postintrarectal administration of TNBS. Five-month-old male *Gatm*^{+/+} ($n = 4$), *Gatm*^{c/+} ($n = 5$), and *Gatm*^{c/c} ($n = 5$) mice were administered 3 mg of TNBS intrarectally, and colons were harvested and measured 4 d posttreatment. * $P < 0.05$; determined by Student's *t* test.

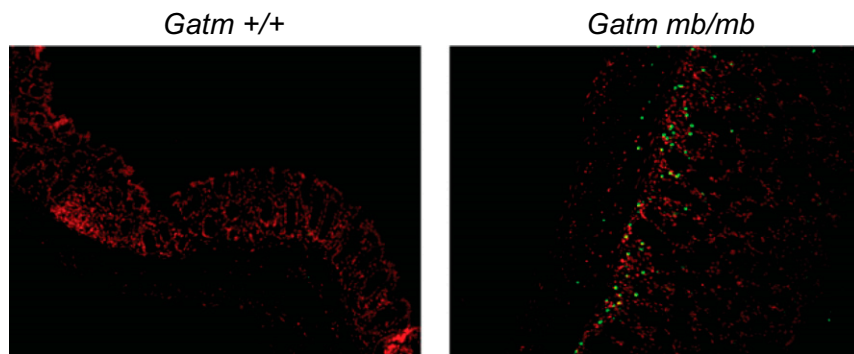


Fig. S3. TUNEL staining of *Gatm*^{mb/mb} colons. Images of TUNEL staining (green) of colon sections from *Gatm*^{+/+} and *Gatm*^{mb/mb} mice on day 6 of 1.5% DSS treatment (10 \times magnification); images are representative of three independent experiments.

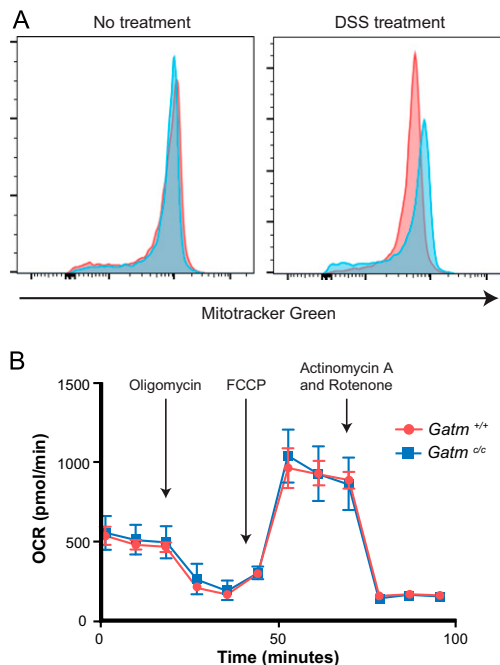


Fig. S4. Mitochondrial mass and functioning in *Gatm*^{c/c} colonocytes. (A) Colonocytes from untreated and 3 d DSS-treated *Gatm*^{c/c} (blue) vs. *Gatm*^{+/+} littermates (red) were harvested and stained with MitoTracker Green. Data are representative of three independent experiments. (B) Mitochondrial respiration of colonic crypts from mice of the indicated genotypes treated for 3 d with 1.5% DSS was tested using Seahorse Flux. Oligomycin, carbonyl cyanide 4-(trifluoromethoxy) phenylhydrazone (FCCP), and a mix of rotenone and antimycin A were serially injected to measure ATP-linked respiration, maximal respiration, and non-mitochondrial respiration, respectively. Data are representative of three independent experiments.